A Reliable Research Partner in Life Science and Medicine

# **MCM3AP Polyclonal Antibody**

catalog number: E-AB-15058

Note: Centrifuge before opening to ensure complete recovery of vial contents.

#### Description

Reactivity Human

Immunogen Recombinant protein of human MCM3AP

Host Rabbit Isotype IgG

**Purification** Affinity purification

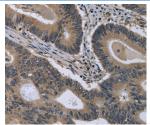
**Buffer** Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.

# **Applications** Recommended Dilution

**WB** 1:500-1:2000 **IHC** 1:50-1:200

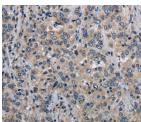
## Data





Western Blot analysis of Hela and 293T cell using MCM3AP Immunohistochemistry of paraffin-embedded Human colon Polyclonal Antibody at dilution of 1:350 cancer using MCM3AP Polyclonal Antibody at dilution of

#### Calculated-MW:218 kDa



Immunohistochemistry of paraffin-embedded Human liver cancer using MCM3AP Polyclonal Antibody at dilution of

1:30

# **Preparation & Storage**

Storage Storage Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.

**Shipping** The product is shipped with ice pack, upon receipt, store it immediately at the

temperature recommended.

## Background

# For Research Use Only

Toll-free: 1-888-852-8623 Web:<u>w w w .elabscience.com</u>

# **Elabscience Bionovation Inc.**



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The minichromosome maintenance protein 3 (MCM3) is one of the MCM proteins essential for the initiation of DNA replication. The protein encoded by this gene is a MCM3 binding protein. It was reported to have phosphorylation-dependent DNA-primase activity, which was up-regulated in antigen immunization induced germinal center. This protein was demonstrated to be an acetyltransferase that acetylates MCM3 and plays a role in DNA replication. The mutagenesis of a nuclear localization signal of MCM3 affects the binding of this protein with MCM3, suggesting that this protein may also facilitate MCM3 nuclear localization. This gene is expressed in the brain or in neuronal tissue. An allelic variant encoding amino acid Lys at 915, instead of conserved Glu, has been identified in patients with mild intellectual disability.

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